# POR gene

cytochrome p450 oxidoreductase

#### **Normal Function**

The *POR* gene provides instructions for making the enzyme cytochrome P450 oxidoreductase. This enzyme is required for the normal functioning of more than 50 enzymes in the cytochrome P450 family. Cytochrome P450 enzymes are involved in the formation (synthesis) and breakdown (metabolism) of various molecules and chemicals within cells.

Cytochrome P450 enzymes are critical for the synthesis of cholesterol and steroid hormones. Cholesterol is a substance that has many essential functions both before and after birth, including roles in the production of steroid hormones and in the formation and growth of bones. Steroid hormones are needed for normal development and reproduction. This group of hormones includes testosterone and estrogen, which are essential for normal sexual development and reproduction; corticosteroids, which are involved in the body's response to stress; and aldosterone, which helps regulate the body's salt and water balance.

Additionally, cytochrome P450 enzymes are involved in the metabolism of ingested substances, such as medications, in the liver. Because cytochrome P450 oxidoreductase helps regulate the activity of these enzymes, researchers suspect that normal variations in the *POR* gene may influence a person's response to particular drugs (drug metabolism).

# **Health Conditions Related to Genetic Changes**

cytochrome P450 oxidoreductase deficiency

More than 50 mutations in the *POR* gene have been found to cause cytochrome P450 oxidoreductase deficiency. This condition causes hormonal changes that can affect the development of the reproductive system, skeleton, and other parts of the body. The disorder affects sexual development before birth and at puberty, and severe cases are also characterized by skeletal abnormalities.

Most of the mutations that cause cytochrome P450 oxidoreductase deficiency change single protein building blocks (amino acids) in cytochrome P450 oxidoreductase. *POR* gene mutations significantly reduce the enzyme's activity, which disrupts the production of steroid hormones. Changes in sex hormones such as testosterone and estrogen lead to problems with sexual development.

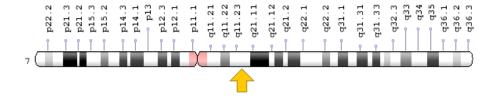
Reduced activity of cytochrome P450 oxidoreductase can also disrupt the production of cholesterol, which likely impairs normal bone formation in severe cases of cytochrome P450 oxidoreductase deficiency. Studies suggest that a molecule called retinoic acid also plays a role in the skeletal abnormalities found in severe cases. The breakdown of retinoic acid requires cytochrome P450 oxidoreductase; if a shortage of cytochrome P450 oxidoreductase prevents retinoic acid from being broken down, the resulting excess of that molecule can stimulate the abnormal growth and fusion of bones.

It is unclear whether mutations in the *POR* gene affect how the liver processes medications. The role of this enzyme in drug metabolism is an active area of research.

#### **Chromosomal Location**

Cytogenetic Location: 7q11.23, which is the long (q) arm of chromosome 7 at position 11.23

Molecular Location: base pairs 75,915,102 to 75,986,855 on chromosome 7 (Homo sapiens Annotation Release 108, GRCh38.p7) (NCBI)



Credit: Genome Decoration Page/NCBI

### Other Names for This Gene

- CPR
- CYPOR
- cytochrome P450 reductase
- FLJ26468
- NADPH-dependent cytochrome P450 reductase
- NCPR HUMAN
- P450 (cytochrome) oxidoreductase
- P450R

# **Additional Information & Resources**

# **Educational Resources**

- Biochemistry (fifth edition, 2002): Steroid hormones https://www.ncbi.nlm.nih.gov/books/NBK22339/#A3657
- Biochemistry (fifth edition, 2002): The Cytochrome P450 System Is Widespread and Performs a Protective Function https://www.ncbi.nlm.nih.gov/books/NBK22339/#A3663
- The Cell: A Molecular Approach (second edition, 2000): Steroid Hormones and the Steroid Receptor Superfamily https://www.ncbi.nlm.nih.gov/books/NBK9924/#A2202

#### GeneReviews

 Cytochrome P450 Oxidoreductase Deficiency https://www.ncbi.nlm.nih.gov/books/NBK1419

# Scientific Articles on PubMed

PubMed

https://www.ncbi.nlm.nih.gov/pubmed?term=%28P450+oxidoreductase%5BTIAB %5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena %5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D

#### OMIM

 CYTOCHROME P450 OXIDOREDUCTASE http://omim.org/entry/124015

#### Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology http://atlasgeneticsoncology.org/Genes/GC\_POR.html
- ClinVar https://www.ncbi.nlm.nih.gov/clinvar?term=POR%5Bgene%5D
- HGNC Gene Symbol Report http://www.genenames.org/cgi-bin/gene\_symbol\_report?q=data/ hgnc\_data.php&hgnc\_id=9208
- NCBI Gene https://www.ncbi.nlm.nih.gov/gene/5447
- UniProt http://www.uniprot.org/uniprot/P16435

# **Sources for This Summary**

- Arlt W, Walker EA, Draper N, Ivison HE, Ride JP, Hammer F, Chalder SM, Borucka-Mankiewicz M, Hauffa BP, Malunowicz EM, Stewart PM, Shackleton CH. Congenital adrenal hyperplasia caused by mutant P450 oxidoreductase and human androgen synthesis: analytical study. Lancet. 2004 Jun 26; 363(9427):2128-35.
  - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15220035
- Flück CE, Tajima T, Pandey AV, Arlt W, Okuhara K, Verge CF, Jabs EW, Mendonça BB, Fujieda K, Miller WL. Mutant P450 oxidoreductase causes disordered steroidogenesis with and without Antley-Bixler syndrome. Nat Genet. 2004 Mar;36(3):228-30. Epub 2004 Feb 1.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/14758361
- Fukami M, Horikawa R, Nagai T, Tanaka T, Naiki Y, Sato N, Okuyama T, Nakai H, Soneda S, Tachibana K, Matsuo N, Sato S, Homma K, Nishimura G, Hasegawa T, Ogata T. Cytochrome P450 oxidoreductase gene mutations and Antley-Bixler syndrome with abnormal genitalia and/or impaired steroidogenesis: molecular and clinical studies in 10 patients. J Clin Endocrinol Metab. 2005 Jan; 90(1):414-26. Epub 2004 Oct 13.
  - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15483095
- Hart SN, Zhong XB. P450 oxidoreductase: genetic polymorphisms and implications for drug metabolism and toxicity. Expert Opin Drug Metab Toxicol. 2008 Apr;4(4):439-52. doi: 10.1517/17425255.4.4.439. Review.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18433346
- Huang N, Agrawal V, Giacomini KM, Miller WL. Genetics of P450 oxidoreductase: sequence variation in 842 individuals of four ethnicities and activities of 15 missense mutations. Proc Natl Acad Sci U S A. 2008 Feb 5;105(5):1733-8. doi: 10.1073/pnas.0711621105. Epub 2008 Jan 29. Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18230729
   Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2234213/
- Huang N, Pandey AV, Agrawal V, Reardon W, Lapunzina PD, Mowat D, Jabs EW, Van Vliet G, Sack J, Flück CE, Miller WL. Diversity and function of mutations in p450 oxidoreductase in patients with Antley-Bixler syndrome and disordered steroidogenesis. Am J Hum Genet. 2005 May;76(5): 729-49. Epub 2005 Mar 25.
   Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/15793702
- Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC1199364/
   Miller WL, Huang N, Agrawal V, Giacomini KM. Genetic variation in human P450 oxidoreductase. Mol Cell Endocrinol. 2009 Mar 5;300(1-2):180-4. doi: 10.1016/j.mce.2008.09.017. Epub 2008 Sep
  - Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/18930113
- Sim SC, Miller WL, Zhong XB, Arlt W, Ogata T, Ding X, Wolf CR, Flück CE, Pandey AV, Henderson CJ, Porter TD, Daly AK, Nebert DW, Ingelman-Sundberg M. Nomenclature for alleles of the cytochrome P450 oxidoreductase gene. Pharmacogenet Genomics. 2009 Jul;19(7):565-6. doi: 10.1097/FPC.0b013e32832af5b7.

Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/19535965
Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2753199/

Reprinted from Genetics Home Reference: https://ghr.nlm.nih.gov/gene/POR

26. Review.

Reviewed: March 2014 Published: March 21, 2017

Lister Hill National Center for Biomedical Communications U.S. National Library of Medicine National Institutes of Health Department of Health & Human Services